AMENDMENT

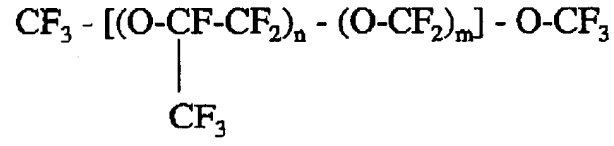
In the claims:

Please cancel claims 14, 20-22, and 24.
Please amend claims 1, 31, 32, and 58 as shown in the following Listing of the Claims.

Listing of Claims:

1. (Currently Amended) A pharmaceutical composition comprising, in addition to one or more pharmacologically active ingredients, wherein the active ingredient is Troxerutine, Nimesulide, a selective CO-2 inhibitor, or a non-steroidal anti-inflammatory drug, wherein said non-steroidal anti-inflammatory drug is Ketoprofen, Diclofenac Sodium, Ibuprofen, Etodolic Acid, Piroxicam, or a combination thereof,

between 0.01 per cent and 60 per cent by weight of a compound of formula I



(I)

with n and m > 18 and < 46 and with a molecular weight between about 600 and about 18,000, in combination with 0.01% to 20% by weight of phosphatidylcholine, for enhancement of active-ingredient absorption.

2. (Canceled)

3. (Previously presented) A pharmaceutical composition according to claim 1 with 0.1 per cent to 30 per cent by weight of the compound of formula I with n and m > 24 and < 36 and with the molecular weight between 1,000 and 4,000.

4. (Previously presented) A pharmaceutical composition according to claim 1, wherein the composition is in a form selected from the group consisting of creams, emulsions, ointments, lotions, foams, gels, aspersion powders, and transdermal formulations.

5-12. (Canceled)

13. (Previously presented) A method for enhancing absorption of a pharmacologically active ingredient, wherein the method comprises topically applying the pharmaceutical composition claimed in Claim 1 to a patient in need thereof, wherein the active ingredient is absorbed through derma, cutis, mucosa, rectum, vagina, or urethra.

14. (Canceled)

15. (Previously presented) A pharmaceutical composition according to Claim 3, wherein the composition is in a form selected from the group consisting of creams, emulsions, ointments, lotions, foams, gels, aspersion powders, and transdermal formulations.

16-17. (Canceled)

18. (Previously presented) The composition according to Claim 1, wherein trans- absorption of the active ingredient is increased by up to more than five times its normal value.

19. (Previously presented) The composition according to Claim 3, wherein trans-absorption of the active ingredient is increased by up to more than five times its normal value.

20. (Canceled)

21. (Canceled)

22. (Canceled)

23. (Previously presented) A method for enhancing absorption of a pharmacologically active ingredient, wherein the method comprises topically applying the pharmaceutical composition claimed in Claim 3 to a patient in need thereof, wherein the active ingredient is absorbed through derma, cutis, mucosa,

rectum, vagina, or urethra.

24. (Canceled)

25. (Previously presented) A pharmaceutical composition as claimed in Claim 1, wherein trans-absorption of the active ingredient is increased by up to more than ten times its normal value.

26. (Previously presented) A pharmaceutical composition as claimed in Claim 1, wherein trans-absorption of the active ingredient is increased by up to more than 20 times its normal value.

27. (Previously presented) A method as claimed in Claim 13, wherein trans-absorption of the active ingredient is increased by up to more than ten times its normal value.

28. (Previously presented) A method as claimed in Claim 13, wherein trans-absorption of the active ingredient is increased by up to more than 20 times its normal value.

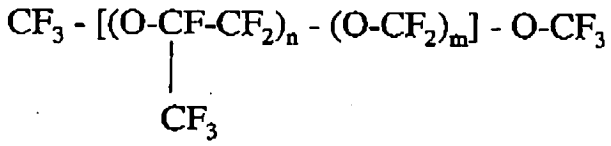
29. (Previously presented) A pharmaceutical composition as claimed in Claim 1, wherein the active ingredient is troxerutine.

30. (Previously presented) A method as claimed in Claim 13, wherein the active ingredient is troxerutine.

31. (Currently amended) A pharmaceutical composition consisting essentially of:

(1) one or more pharmacologically active ingredients, wherein the active ingredient is Troxerutine, Nimesulide, a selective CO-2 inhibitor, or a non-steroidal anti-inflammatory drug, wherein said non-steroidal anti-inflammatory drug is Ketoprofen, Diclofenac Sodium, Ibuprofen, Etodolic Acid, Piroxicam, or a combination thereof;

(2) between about 0.01 per cent and about 60 per cent by weight of a compound of formula I



(I)

wherein n and m are each greater than 18 and are each less than 46 and wherein the compound of the formula I has a molecular weight between about 600 and about 8,000;

(3) phosphatidylcholine;

(4) optionally tocopherol acetate;

(5) optionally polyacrylamide, C_{13}-C_{14} isoparaffin, and laureth-7;

(6) optionally methyl-p-hydroxybenzoate;

(7) optionally propyl-p-hydroxybenzoate;

(8) optionally phenoxyethanol;

(9) optionally nor-chenodeoxycolic acid;

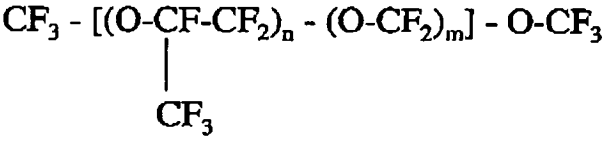
(10) optionally transcutol; and

(11) optionally water.

32. (Currently amended) A pharmaceutical composition consisting essentially of:

(1) one or more pharmacologically active ingredients, wherein the active ingredient is Troxerutine, Nimesulide, a selective CO-2 inhibitor, or a non-steroidal anti-inflammatory drug, wherein said non-steroidal anti-inflammatory drug is Ketoprofen, Diclofenac Sodium, Ibuprofen, Etodolic Acid, Piroxicam, or a combination thereof,;

(2) between about 0.01 per cent and about 60 per cent by weight of a compound of formula I



(I)

wherein n and m are each greater than 18 and are each less than 46 and wherein the compound of the formula I has a molecular weight between about 600 and about 8,000;

(3) phosphatidylcholine;

(4) optionally tocopherol acetate;

(5) optionally polyacrylamide, C_{13}-C_{14} isoparaffin, and laureth-7;

(6) optionally methyl-p-hydroxybenzoate;
(7) optionally propyl-p-hydroxybenzoate;
(8) optionally phenoxyethanol;
(9) optionally nor-chenodeoxycolic acid;
(10) optionally transcutol;
(11) optionally lactic acid;
(12) optionally ethyl alcohol; and
(13) optionally water.

33. (Previously presented) A pharmaceutical composition as claimed in Claim 31, wherein the active ingredient is troxerutine.

34. (Previously presented) A pharmaceutical composition according to claim 31, wherein the phosphatidlycholine constitutes 0.01 per cent to 10 per cent by weight of the pharmaceutical composition, and wherein the compound of the formula I has a molecular weight between 1,000 and about 4,000 with n and m each greater than 24 and each less than 36.

35. (Previously presented) A pharmaceutical composition according to claim 33, wherein the phosphatidlycholine constitutes 0.01 per cent to 10 per cent by weight of the pharmaceutical composition, and wherein the compound of the formula I has a molecular weight between 1,000 and about 4,000 with n and m each greater than 24 and each less than 36.

36. (Previously presented) A pharmaceutical composition as claimed in Claim 32, wherein the active ingredient is troxerutine.

37. (Previously presented) A pharmaceutical composition according to claim 32, wherein the phosphatidlycholine constitutes 0.01 per cent to 10 per cent by weight of the pharmaceutical composition, and wherein the compound of the formula I has a molecular weight between 1,000 and about 4,000 with n and m each greater than 24 and each less than 36.

38. (Previously presented) A pharmaceutical composition according to claim 36, wherein the phosphatidlycholine constitutes 0.01 per cent to 10 per cent by weight of the pharmaceutical composition, and wherein the compound of the

formula I has a molecular weight between 1,000 and about 4,000 with n and m each greater than 24 and each less than 36.

39. (Previously presented) The method according to Claim 13, wherein the active ingredient is Troxerutine, Nimesulide, Ketopropfen, Etodolic Acid, or a combination thereof.

40. (Previously presented) The pharmaceutical composition as claimed in Claim 1, wherein the active ingredient is Troxerutine, Nimesulide, Ketopropfen, Etodolic Acid, or a combination thereof.

41. (Previously presented) The pharmaceutical composition as claimed in Claim 31, wherein the active ingredient is Troxerutine, Nimesulide, Ketopropfen, Etodolic Acid, or a combination thereof.

42. (Previously presented) The pharmaceutical composition as claimed in Claim 32, wherein the active ingredient is Troxerutine, Nimesulide, Ketopropfen, Etodolic Acid, or a combination thereof.

43. (Previously presented) The pharmaceutical composition as claimed in Claim 1, wherein phosphatidylcholine is 0.01% to 10% by weight of the pharmaceutical composition.

44 - 57 (Canceled)

58. (Currently amended) The pharmaceutical composition as claimed in claim 31, wherein the composition consists essentially of the one or more active ingredients, wherein the active ingredient is Troxerutine, Nimesulide, a selective CO-2 inhibitor, or a non-steroidal anti-inflammatory drug, wherein said non-steroidal anti-inflammatory drug is Ketoprofen, Diclofenac Sodium, Ibuprofen, Etodolic Acid, Piroxicam, or a combination thereof, the compound formula I, the phosphatidylcholine, and optionally the water.

59. (Previously presented) A method according to Claim 13, wherein trans-absorption of the active ingredient is increased by up to more than five times its normal value.